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Aftovaxpur DOE (WD) (02) O1 BFS

Not authorised

- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated
- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated

Product identification

Medicine name:
Aftovaxpur DOE (WD) (02) O1 BFS

Active substance:
Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated

Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated

Target species:
Cattle
Sheep
Pig
Sheep
Pig
Cattle

Route of administration:
Intramuscular use
Subcutaneous use
Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated

Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated

Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Not applicable. 0 day Zero days
- **Sheep**
  - Not applicable. 0 day Zero days
- **Pig**
  - Not applicable. 0 day Zero days

Subcutaneous use:

- **Cattle**
  - Not applicable. 0 day Zero days
- **Sheep**

- Not applicable. 0 day Zero days

- **Pig**
  - Not applicable. 0 day Zero days

Subcutaneous use:

- **Sheep**
  - Not applicable. 0 day Zero days

- **Pig**
  - Not applicable. 0 day Zero days

- **Cattle**
  - Not applicable. 0 day Zero days

Intramuscular use:

- **Cattle**
  - Not applicable. 0 day Zero days

- **Sheep**
  - Not applicable. 0 day Zero days

- **Pig**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AA04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Surrendered

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
15/07/2013

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
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Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
15/07/2013

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Documents

Combined File of all Documents

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Published on: 14/03/2026
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